

ADVT. NO. 09-03
Website Ref: RollingAdvtSept2104

- Project titled:** Establishment of National Repository for cGMP complaint microbial Cell bank of Biopharmaceutical Relevance (NRGCBIO)

Project Code: GAP-0195

Position: GMP & Regulatory Compliance Consultant= 01

Essential Qualification: Should have at least Graduate degree in any engineering & technology/Biochemical Engineering/Biological Sciences. Having Experience in the field of cGMP operations, Biopharmaceuticals development (Biotherapeutics, Vaccine, Biosimilar, immunotherapeutics etc), quality management systems, GMP facility design, setup, facility validation & engineering, regulatory affairs in the field of Biopharmaceuticals in line with national and international norms as evidenced by working experience certificates. Should have experience to review GMP facility setup documents, facility validation, compliance etc.

Three letters of recommendation from senior level Industry leaders/Scientists/Faculty is required.

Desirable: 10-15 years of work experience in projects in Biopharmaceutical facility/labs/biotech industry/reputed engineering consultancy firms. Must have through knowledge of Indian, ICH, USFDA, EU, guidelines essential for setting up and qualification of GMP compliant facility. Experience in Biosafety & Biocontainment Requirements, industrial safety, environmental compliance in Biopharmaceuticals R&D. Post-graduate/Ph.D. degree is desirable.

Monthly Emoluments: Rs. 1, 00000 (One Lakh) /Month Consolidated

Selection procedure: The engagement will be purely on contract basis. Selections will be based on walk in interview in response to this advertisement, on the basis of experience and qualification of applicants. Consultants will be selected on the basis of interview.

NOTE: Last Date of Receipt of Applications – 15th October, 2021.

CONTROLLER OF ADMINISTRATION